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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/525,630	10/24/2005	Alex Zhu	26471-501 NATL	7794
30623 7590 01/14/2008 MINTZ, LEVIN, COHN, FERRIS, GLOVSKY AND POPEO, P.C.			EXAMINER	
			HOLT, ANDRIAE M	
ONE FINANC BOSTON, MA			ART UNIT	PAPER NUMBER
BOSTON, MA	. 02111		1616	
			MAIL DATE	DELIVERY MODE
			01/14/2008	PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

	Application No.	Applicant(s)				
Office Action Summary	10/525,630	ZHU, ALEX				
Office Action Summary	Examiner	Art Unit				
	Andriae M. Holt	1616				
The MAILING DATE of this communication app Period for Reply	ears on the cover sneet with the c	orrespondence address				
A SHORTENED STATUTORY PERIOD FOR REPLY WHICHEVER IS LONGER, FROM THE MAILING DA - Extensions of time may be available under the provisions of 37 CFR 1.13 after SIX (6) MONTHS from the mailing date of this communication. - If NO period for reply is specified above, the maximum statutory period w - Failure to reply within the set or extended period for reply will, by statute, Any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b).	ATE OF THIS COMMUNICATION 16(a). In no event, however, may a reply be tim rill apply and will expire SIX (6) MONTHS from cause the application to become ABANDONEI	I. ely filed the mailing date of this communication. O (35 U.S.C. § 133).				
Status						
1) Responsive to communication(s) filed on 24 Oc	<u>ctober 2005</u> .					
2a) This action is FINAL . 2b) This	This action is FINAL . 2b) This action is non-final.					
·	Since this application is in condition for allowance except for formal matters, prosecution as to the merits is					
closed in accordance with the practice under Ex parte Quayle, 1935 C.D. 11, 453 O.G. 213.						
Disposition of Claims						
4) ⊠ Claim(s) <u>1-86</u> is/are pending in the application. 4a) Of the above claim(s) is/are withdraw 5) □ Claim(s) is/are allowed. 6) □ Claim(s) is/are rejected. 7) □ Claim(s) is/are objected to. 8) ⊠ Claim(s) <u>1-86</u> are subject to restriction and/or expressions.	vn from consideration.					
Application Papers						
9) The specification is objected to by the Examine 10) The drawing(s) filed onis/ are: a) acce Applicant may not request that any objection to the Replacement drawing sheet(s) including the correct 11) The oath or declaration is objected to by the Ex	epted or b) objected to by the Edrawing(s) be held in abeyance. See ion is required if the drawing(s) is obj	e 37 CFR 1.85(a). ected to. See 37 CFR 1.121(d).				
Priority under 35 U.S.C. § 119						
12) Acknowledgment is made of a claim for foreign a) All b) Some * c) None of: 1. Certified copies of the priority documents 2. Certified copies of the priority documents 3. Copies of the certified copies of the prior application from the International Bureau * See the attached detailed Office action for a list	s have been received. s have been received in Applicati ity documents have been receive u (PCT Rule 17.2(a)).	on No ed in this National Stage				
Attachment(s) 1) Notice of References Cited (PTO-892) 2) Notice of Draftsperson's Patent Drawing Review (PTO-948) 3) Information Disclosure Statement(s) (PTO/SB/08) Paper No(s)/Mail Date	4) Interview Summary Paper No(s)/Mail Da 5) Notice of Informal P 6) Other:	ate				

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DETAILED ACTION

Election/Restrictions

Restriction is required under 35 U.S.C. 121 and 372.

This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1.

In accordance with 37 CFR 1.499, applicant is required, in reply to this action, to elect a single invention to which the claims must be restricted.

Group I, claim(s) 1-10 and 43-47, drawn to a method and a pharmaceutical composition for localized chemodenervation comprising administering a composition comprising an antibiotic and a pharmaceutically acceptable carrier.

Group II, claim(s) 11-16 and 60-67, drawn to a method and a pharmaceutical composition for localized chemodenervation comprising administering a composition comprising a muscle relaxant and a pharmaceutically acceptable carrier.

Group III, claim(s) 17-19 and 48-52, drawn to a method and a pharmaceutical composition for localized chemodenervation comprising administering a composition comprising a plant extract and a pharmaceutically acceptable carrier.

Group IV, claim(s) 20-25 and 69-73, drawn to a method and a pharmaceutical composition of treating involuntary facial muscle spasms.

Group V, claim(s) 26-30 and 75-59, drawn to a method and a pharmaceutical composition for reducing facial wrinkles.

Group VI, claim(s) 31-36 and 81-85, drawn to a method and a pharmaceutical composition for treating or reducing neuropathic pain.

Group VII, claim(s) 37-42, drawn to a pharmaceutical composition for topical administration for localized chemodenervation comprising a polymyxin and a magnesium salt.

Group VIII, claim(s) 53-55, drawn to a pharmaceutical composition for topical administration for localized chemodenervation comprising tetracycline in a pharmaceutically acceptable medium.

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Group IX, claim(s) 56-59, drawn to a pharmaceutical composition for topical administration for localized chemodenervation comprising lincosamide in a pharmaceutically acceptable medium.

Group X, claim(s) 68, drawn to a kit for treatment of localized chemodenervation.

Group XI, claim(s) 74, drawn to a kit to treat involuntary muscle spasms.

Group XII, claim(s) 80, drawn to a kit to reduce facial wrinkles.

Group XIII, claim(s) 86, drawn to a kit to treat or reduce neuropathic pain.

The inventions listed as Groups I - XIII do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons: There is no special technical feature. The groups are to various inventions that lack a special technical feature. Group I is drawn to a method for localized chemodenervation using an antibiotic, whereas, group II is drawn to a method for localized chemodenervation using a muscle relaxant. Each group is to treat a localized chemodenervation, but with a different type of drug. Group I is drawn to a method and pharmaceutical composition for localized chemodenervation using an antibiotic, whereas, group V is a method and pharmaceutical composition to reduce facial wrinkles, the groups lack a special technical feature, as each group is a composition or a method to treat various ailments or symptoms, ranging from facial wrinkles to neuropathic pain.

The examiner has required restriction between product and process claims.

Where applicant elects claims directed to the product, and the product claims are subsequently found allowable, withdrawn process claims that depend from or otherwise

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require all the limitations of the allowable product claim will be considered for rejoinder.

All claims directed to a nonelected process invention must require all the limitations of an allowable product claim for that process invention to be rejoined.

In the event of rejoinder, the requirement for restriction between the product claims and the rejoined process claims will be withdrawn, and the rejoined process claims will be fully examined for patentability in accordance with 37 CFR 1.104. Thus, to be allowable, the rejoined claims must meet all criteria for patentability including the requirements of 35 U.S.C. 101, 102, 103 and 112. Until all claims to the elected product are found allowable, an otherwise proper restriction requirement between product claims and process claims may be maintained. Withdrawn process claims that are not commensurate in scope with an allowable product claim will not be rejoined. See MPEP § 821.04(b). Additionally, in order to retain the right to rejoinder in accordance with the above policy, applicant is advised that the process claims should be amended during prosecution to require the limitations of the product claims. Failure to do so may result in a loss of the right to rejoinder. Further, note that the prohibition against double patenting rejections of 35 U.S.C. 121 does not apply where the restriction requirement is withdrawn by the examiner before the patent issues. See MPEP § 804.01.

Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a request under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

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Applicant is advised that the reply to this requirement to be complete must include (i) an election of a species or invention to be examined even though the requirement be traversed (37 CFR 1.143) and (ii) identification of the claims encompassing the elected invention.

The election of an invention or species may be made with or without traverse. To reserve a right to petition, the election must be made with traverse. If the reply does not distinctly and specifically point out supposed errors in the restriction requirement, the election shall be treated as an election without traverse.

Should applicant traverse on the ground that the inventions or species are not patentably distinct, applicant should submit evidence or identify such evidence now of record showing the inventions or species to be obvious variants or clearly admit on the record that this is the case. In either instance, if the examiner finds one of the inventions unpatentable over the prior art, the evidence or admission may be used in a rejection under 35 U.S.C.103 (a) of the other invention.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Andriae M. Holt whose telephone number is 571-272-9328. The examiner can normally be reached on 7:00 am-4:00 pm.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Richter Johann can be reached on 571-272-0646. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

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Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

Andriae M. Holt **Patent Examiner** Art Unit 1616

Johann R. Richter

Supervisory Patent Examiner

Art Unit 1616